Data Validation Specialist

TRIO (Translational Research in Oncology) is a not-for-profit clinical research organization that is dedicated to advancing translational cancer research by pursuing forward innovative and targeted therapeutic concepts in the clinical trial setting. We are committed to providing treatments of the future to the world of today.

TRIO’s head office is in Edmonton, Alberta, Canada, although the organization has operations all throughout Canada, the USA, France and Uruguay. TRIO is looking for a Data Validation Specialist to join our Edmonton office.

Responsibilities:
- Review clinical data collected in the patient electronic Case Report Form (eCRF) in order to ensure consistency and accuracy of the data.
- Generate queries to the sites where inconsistencies in the data or missing data has occurred.
- Review answers to the queries, provided by the sites, in order to ensure adequate resolution of the issues found.
- Track the progress of the data cleaning activity compared to the study plan in order to adequately plan trial analyses.
- Track protocol deviations identified during the data cleaning process.
- Participate in the review of other study documents (e.g. guidelines, specifications, newsletters, manuals and reports) associated with data validation activities.

Qualifications:
- Medical background is preferred.
- Bachelor’s degree, in a science/health related field.
- Minimum one-year experience in Clinical Research, or medical field.
- Demonstrate high attention to detail, accuracy and thoroughness.
- Fluent in English, written and verbal.
- Work well under pressure to meet project deadlines.
- Intermediate proficiency in MS Office and other software applications.

If this sounds like the opportunity you have been seeking please forward a cover letter with your CV by email, quoting 19-25 DVS in the subject line to: human.resources@trioncology.org.

We thank all candidates for their interest; only those selected for an interview will be contacted.